Scientists Call for Global Moratorium on mRNA Vaccines, Immediate Removal from Childhood Schedule

A new open letter published in the journal Science is the first pier reviewed paper to call for a global moratorium on the COVID-19 mRNA vaccines. The authors say that the vaccines have failed to live up to their promises, and that the benefits of the mRNA technology for other vaccine platforms should not have been overshadowed as it has.

By **Brenda Baletti, Ph.D.**

Ultra-clip, Mike A. Subtitle to the Skeptic's Top Many of the Q's, **It's free.**

Governments should endorse a global moratorium on mRNA vaccines and all questions about their safety have been thoroughly investigated, according to the authors of a new, peer reviewed article on the COVID-19 vaccine trials and the mRNA vaccine platform campaign published this week in the journal Science Medical Science.

Curious is a web-based peer-reviewed open-access general medical journal covering all medical sciences.

The authors surveyed published research on the pharmaceutical companies' COVID-19 vaccine trials, including Pfizer's BNT162b2 and Moderna's mRNA-1273, and found numerous methodological errors in the childhood immunization schedule.

After the first-reported vaccine event claimed they were 95% effective in preventing COVID-19, serious problems with method, execution and reporting in the trial vaccine public, which this paper reviewed in detail.

Evidence also shows the products never underwent adequate safety and effectiveness assessments before they were rolled out to billions of people and a significant number of serious injuries and deaths have been reported.

The article was published Wednesday, September 29th, 2021, by Science Medical Science.

It is the first peer-reviewed study to call for a moratorium on the COVID-19 mRNA vaccines, and the mRNA platform.

"After a proper assessment of the safety and efficacy criteria, we made haste — upon which the emergency use authorization (EUA) and ultimate full authorization were granted by the FDA — despite the limited data from the experimental trials.

They concluded, "Federal agency approval of the COVID-19 mRNA injectable products on a blanket coverage population-wide basis had no support from an adequate assessment of safety and efficacy, nor from a comprehensive consideration of risks versus benefits."

They also called for the vaccines to be immediately removed from the childhood immunization schedule and for the cessation of the boosters.

"It is unethical and unconscionable to administer an experimental vaccine to a child who is not at risk from the disease it is meant to prevent," they wrote, and established 2.2% risk of permanent heart damage based on the best prospective data available. They wrote:

Finally, the authors called for a full investigation into misconduct by the pharmaceutical companies and the regulatory agencies.

It is the first peer-reviewed study to call for a moratorium on the COVID-19 mRNA vaccines, and the mRNA platform.

The authors suggested that investigation should include:

- Whether safety and efficacy criteria were made forgotten;
- Whether the emergency use authorizations (EUA) and ultimate full authorizations were granted by the FDA — despite the limited data from the experimental trials;
- Whether safe vs effective; and
- Whether the COVID-19 mRNA injectable products were rushed into authorization.

According to McCullough, mRNA should never have been authorized for human use.

Lead author Ward Stat Defender, "Our claim is that any risk-benefit analysis must consider how the risk presented to human health by the mRNA technology itself is related mortality is offset by the potential increase in vaccine-induced mortality."

Here are six takeaways from the review:

1. The COVID-19 vaccines' are rushed-to-market gene therapies that were rushed through the regulatory process in a historically unprecedented manner

Before the seven-month authorization process for the mRNA vaccines, no vaccine had ever gone to market without undergoing testing of at least four years, with typical timelines averaging 10 years.

To speed the process, the companies skipped preclinical studies of potential toxicity, genotoxicity, carcinogenicity, and studies on impact on fertility, standard for identifying longer-term adverse effects and the established 10-year period for monitoring for long-term effects such as cancer and autoimmune disorders, the authors wrote.

The trials prioritized documenting effective symptom reduction over SAE and death, "which is inappropriate for assessing the risk-benefit balance of gene therapy products and gene therapy products rehabilitated as vaccines and never given the first-line-zone use for use against a viral disease."

However, the gene therapies' components have not been thoroughly evaluated for safety for use as vaccines.

There is an unrecognized and major concern that the mRNA could transform body cells into viral protein factories — with no off-switch — that produce the spike protein and disseminate pieces of it throughout the circulatory system, the authors wrote.

The spike protein in the vaccine, the authors said, is associated with more severe immunopathology and other dys-than the spike protein in the viral particle.

The authors suggested that massive government investment in mRNA technology, including hundreds of billions by federal agencies and tens of billions in Big Tobacco, "U.S. federal agencies and tens of billions in Big Tobacco, "U.S. federal agencies and tens of billions in Big Tobacco" toward successful outcomes for the regimentation trial."

The financial incentives along with political pressures to deliver a rapid solution likely influenced a series of flawed decisions that compromised the integrity of the clinical trials and eclipsed serious scientific concerns about risks with the technology, the added.

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2. Steps were taken in trials to overestimate vaccine efficacy

Because the trials were designed to assess whether the mRNA vaccine reduced symptoms, they did not measure whether the vaccines prevented severe disease and death. Yet the vaccine makers repeatedly claimed they did.

The large randomized double-blind placebo-controlled trials have been characterized as reductions in SARS-CoV-2 transmission, hospitalization, or death," the authors wrote.

Additionally, the number of people who contracted clinical COVID-19 in both the placebo and intervention groups was low, and the statistical significance, or broad-sweeping conclusions with regard to COVID-19 morbidity and mortality."

Pfizer's 36 % efficacy claims were not large enough to be significant, the authors wrote. Moderna's 94.1% efficacy claim was based on a single dose, and in the placebo recipients died from COVID-19 in the 1-to-two-month, only one placebo death was attributed to COVID-19.

There was also a much larger percentage of "suspected COVID-19 cases" in both groups, with participants showing COVID-19 symptoms but a negative PCR test. When factoring in these cases, measures of vaccine efficacy fell to 19%.

The trial subject pool was comprised of largely young and healthy individuals. Efficacy was not established in the elderly population, which is the population most vaccinated — and most vulnerable to severe disease and death.

Findings from analyses of data from the Pfizer trials can be interpreted as showing the vaccines made "no significant difference" in reducing all-cause mortality, the authors said, and a significant increase was observed in the non-COVID-19 deaths.

Even the six-month grace-marketing data Pfizer presented to the U.S. Food and Drug Administration (FDA) showed no reduction in all-cause mortality, the authors wrote.

The authors maintained that data adjusting the analysis of deaths to better account for the fact that Pfizer unblinded the study people from the placebo group "calls into question the safety of the vaccine." The 2.2% mortality rate and 10.95% that the unvaccinated group (19,965 people) by the end was a conservative estimate.

One of the most glaring errors with the registrational trials, they noted, was that they exclusively focused on measuring risk reduction — the ratio of COVID-19 cases in the placebo group to the vaccine group — "while failing to report on measuring absolute risk reduction, which is the likelihood someone will get COVID-19 symptoms relative to people in the population at large.

According to FDA guidelines, vaccinology for both approaches is crucial to avoid the misguided use of pharmaceutical products — but the data were omitted, leading to the overestimation of an intervention's clinical utility.

While both vaccines touted an approximately 95% risk reduction figure as their efficacy, the absolute risk reductions for the Pfizer and Moderna vaccines were 0.7% and 1.1% respectively.

"A substantial number of individuals would need to be injected in order to prevent a single mild-to-moderate case of COVID-19," the authors wrote.

As an example, using a conservative estimate that 119 people would need to be vaccinated to prevent infection, and assuming that COVID-19 had a 0.3% infection fatality rate, they wrote that approximately 52,500 vaccinations would be necessary to prevent a single COVID-19 fatality.

However, "Given clear misconduct and data-integrity problems ... the true benefit may be even lower than estimated."

Any duly vetted, one would want to assess the benefit along with risks — which this estimate to be 21 deaths per 100,000 doses of Pfizer. However, using the most conservative estimates, "it is therefore unclear if the vaccines do more deaths caused by the modified mRNA injections."

3. Numbers of SAEs in the trials and post-rollout reporting are well-documented, despite claims to the contrary

Both Pfizer and Moderna found about 132 SAEs per 100,000 vaccine recipients, or one SAE for every 800 vaccines. However, because the trials were not designed to identify rare adverse events, the authors note, even higher proportions of SAEs would be expected in the general population.

The Chairman et al. reanalysis of the Pfizer trial data found a significant 36% increase in SAEs in the vaccine recipients and many life-threatening autoimmune events in the vaccinated participants.

Official SAEs for other vaccines average around only 1.2 per million. Chairman et al. determined 1,200 SAEs per million vaccinations, exceeding the benchmark by "at least 600-fold."

After the vaccine rollout, analyses of the two large drug-safety reporting systems in the U.S. and Europe identified signals for myocardial infarction, pulmonary embolism, thrombosis, and Bell's palsy, among other serious adverse events associated with both mRNA vaccines, along with tumors, cancers, and millions of VAIs have reported to those systems.

Another study by Skidmore et al. examined the total number of fatalities from the vaccines in 2021 alone was 289,786. Autopsy studies have also provided evidence of vaccine-induced SAEs, including myocarditis, which is the leading cause of death in athletes, reported the authors.

In multiple autopsy studies, German pathologist Arne Burkhardt documented the presence of vaccine-antibodies spike proteins in blood vessels, organs, and brain tissues. This research helps to explain documented vaccine-induced fatalities affecting the nervous, immune, reproductive and other systems.

The Pfizer data also showed an overwhelming number of adverse effects, including 1,223 deaths and 42,086 serious adverse events (SAEs), documented approximately 1.6 million AEs affecting nearly 40% of trial participants, and one-third of them were classified as severe.

In the U.K., Yellow Card, Pfizer's reports found a nearly 4-fold increase (OR 3.7; 95%CI 1.55-13.2; p = 0.03) in serious cardiac events, including myocarditis, pericarditis, and arrhythmias, in the vaccine group. Moderna's Spikevax (Moderna's mRNA-1273) vaccine was acknowledged as "associated with a risk for safety signal.

"The serious adverse events are all well documented," Wood said. "Yet it's surprising to see so many in the medical field continue to ignore or dismiss them, and to be taken half of the equation when considering the risk-benefit ratio."

4. The failure to appropriately test for safety and toxicity poses serious problems.

Researchers have raised concerns that the mRNA technology is inherently unstable and difficult to create, which leads to batch variability and contamination linked to different rates of safety.

Recent findings by Mörtenson et al. that found Pfizer's mRNA vaccines are contaminated with plasmid DNA that shouldn't be present — and aren't present in the vaccines used in the trials — raising serious safety issues.

That's because "Viruses, 1," used in the trials to generate the vaccines involved in the trials to generate the vaccines, "and the vaccines are contaminated with DNA fragments that violate the manufacturing process for mRNA vaccines," Pfizer's "Viruses, 1," which involves using E. coli bacteria to express the proteins.

Removing plasmids E. coli, can result in residual plasmids in the vaccines and the effects of their presence is unknown.

Mörtenson's work also revealed the presence of DNA from simian virus 40 (SV40), an oncogenic DNA virus originally isolated in 1960 from contaminated polio vaccines administered to millions of people. It is also present in many laboratory animals, raising other safety concerns.

Researchers from Cambridge published a paper in Nature in December 2021, which found that mRNA vaccines activated the multiple mRNA executioners in the spike protein in COVID-19 immunizations that causes the machinery that produces the spike protein to run out of control, leading to cell death.

This process causes a "translation" that can result in the production of harmful proteins in addition to the spikes. These proteins, which developers either failed to look for or did not report in regulatory submissions, could potentially increase the vaccine-induced adverse reactions.

5. There are many different possible biological mechanisms that cause AEs and vaccine failure.

The review points readers to a series of papers that explain a number of different theories to explain how high numbers of AEs from the COVID-19 vaccines.

"The mechanisms of molecular mimicry, antigen cross-reactivity, pathogenic priming, viral reactivation, immune exhaustion, and other factors related to immune dysregulation are important to consider in the development and pathogenesis of malignant and autoimmune diseases," they wrote. And these mechanisms of immune activation are distinct from the body's response to a viral infection.

They also note the toxic effects of the primary adjuvants, PEG, and of the spike protein itself.

They close their analysis of the vaccines with a complex explanation for the "unprecedented" number of AEs reported to the Vaccine Adverse Event Reporting system, naturally-immunity through infection. They explain the mechanisms for vaccine failure and problems generated by the ability for the mRNA vaccines to perpetuate the emergence of new variants.

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Brenda Baletti, Ph.D.

Brenda Baletti, Ph.D. is a reporter for The Defender. She is an avid and eager student of science, nature and health, and has a passion for writing and helping others. She received her Ph.D. in Biochemistry and Molecular Biology from the University of Miami and has worked in both the University of Texas & Stanford.

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comply and fit in. A big piece of the training is how to be socialized into the trade. I used to say the same about going to college when people worried about not committing to a particular major or field to go into. It is 4 yrs of learning how to deal on a particular level in our society. Medical people learn to function on a certain level and get kudos only when they follow the rules. Years of this training, on top of whatever they brought with them from younger years results in producing good and dutiful clones. Remember even at the beginning of covid whenever a doctor or nurse raised questions based on their observation they were rudely shut down, dismissed or isolated. Like getting spanked and sent to the corner until they could behave themself.

4 0 Reply Share

Paul Hastilow
13 days ago

Two Major factors are apparent in this issue.
Financial Incentives, (warp speed Vaxxines by Big pHarma) Billions in Sales.
Political Pressure.....! To find a Remedy..!
Even though there never was any Support for "Honest Assessments" on evidence about "Adverse Reactions" due to "Haphazard" reporting from Federal Agencies, (FDA, CDC, WHO, NIH.)
Fauci and His Cohorts Jead the Charge into "Stupidity". Masks, Lockdowns. (Flatten the Curve). Social Distancing, Infecting your Granny??? Closing Schools. The use of Remdesivir (Repurposed from AIDS), use of Ventilators, (Guaranteed Death).
It was All about "Money" & "Control".... We were/are the Experiment...!
And the "Scientist Dm" don't like us proving them Wrong on so many issues.
They hide behind "Redacted Documents", and when Questioned on the subject,
The Answer is "I don't Recall".... Lies, Lies, Lies.
Give All concerned a "Polygraph Test" or Prison time for Contempt of Court, until their Memory comes back.!
So Sad to see No One is Held accountable for their "Crimes against Humanity.".!

11 0 Reply Share

Double_Up
13 days ago

With turbo cancers, deaths from heart damage, deaths and disorders from neurological damage, what else can the vaxxes do?

11 0 Reply Share

CastleConnell → Double_Up
13 days ago

Mad cow

4 0 Reply Share

el Gallinazo
13 days ago edited

The DOD purchased all the clotshots as **countermeasures** under the Prep Act. They were never legally pharmaceuticals. The Christian Drosten rt-PCR test was a total fraud. It never had a validation study, as required for a chemical based diagnostic test. A Magic Eight Ball would have performed better in a validation study. It was necessary as there never was a SARS-CoV-2 virus so the "pandemic" had to be artificially and exponentially boosted. We finally got a symptomless, lethal disease. A miracle of modern medicine. Kari Mullis died a few months before the start of the scamdemic. He would have been a HUGE thorn in the side of the parasite class regarding their operation. Since the clotshot was legally a defensive weapon system purchased and administered by the DOD, the "trials" were Stalinesque show trials (pun intended) and had no legal standing. Pfizer's defense against civil charges of fraud was that the government, who paid the bill, was in on it, so no one was defrauded. The scamdemic had two primary objectives as it was rolled out by the parasite class:
1) Kill the global economy.
2) Depopulation. I suspect that all or most of the children who got the clotshots when they reach puberty will be discovered to be infertile. Women who got it in the first and second trimesters have had statistically terrible issues with bringing their babies to a healthy term. For everyone who took it (which includes my 50 over my vociferous screams), you have been gaslighted. Even Ingrid Bergman figured it out by the end of the movie.

The real story is being unfolded by Sasha Latypova for people who are interested in the truth.

8 0 Reply Share

George → el Gallinazo
13 days ago

Yep, Sasha Latypova and Katherine Watts have miraculously unraveled the complex web of government laws and rules that deceived most of us. Se their Substack articles. Similar situations have been uncovered in numerous countries including, Canada, Australia, UK, New Zealand, the EU... It's a global operation that was developed over decades! The criminals behind this are not Big Pharma, they just played along. It's DoD, HHS...

8 0 Reply Share

Lionheart → el Gallinazo
12 days ago edited

Thanks for this information, I will look up the ladies mentioned. I recall Clayton & Natalie Morris from Redacted on Youtube few months ago uncovering and discussing official DOD documentation preparing this 'evil' clot shot agenda more than a decade ago. You're right, Big Pharma and even Blackrock (Blackrock Owns Pfizer) just played along, they are not the biggest threat, Unbelievable!!!

3 0 Reply Share

Voyt Regnal
13 days ago

Dream on. It will not happen. Meanwhile avoid these poison death shots for yourself and kids. Create/obtain fake certification if necessary...I predict this will be easier every year.

10 1 Reply Share

CastleConnell → Voyt Regnal
13 days ago

I saw that medical staff who created fake vaccine cards are being prosecuted. I don't recommend that strategy it can come back and cause big legal problems.

2 4 Reply Share

Rob DF → CastleConnell
13 days ago

I didn't take the VAX and couldn't travel for almost 2 years... next time the gloves come off.

6 0 Reply Share

A. Miller
13 days ago

Yeah....but talk about a day late and thousands of deaths and injuries too late...these people are part of the problem.

5 0 Reply Share

tmlogan → A. Miller
13 days ago

If you're speaking of the brave scientists that published this paper, they have been speaking out from the very beginning, not only about the jabs but about all the Covid fear mongering & hospital protocols, and they've been maligned every step of the way. But they didn't quit & now they managed to do this in depth analysis & publish this review, which is quite amazing. They have been part of the problem, and have paid dearly but haven't quit. Unless you've also been willing to lay your career & reputation on the line, you've got nothing to say against their efforts God Bless Them!!!.

10 0 Reply Share

el Gallinazo
13 days ago

Yes, just like your typical mafia hitman gets "bamboozled" by the extra money.

3 0 Reply Share

JSJ
12 days ago

The majority of all degree holders including doctors is in the 115 to 120 range just above average. 80% of the world's population isn't actually capable of actually thinking; They can tell you how they feel about something but if you ask them what they think about it, they don't know what to say. The ability to perform complex Routines on command is mistaken for intelligence. The core of an average Doctor's ego is probably the praise and money they receive for being smart, special, and important.

4 0 Reply Share

Matt
12 days ago

Look at the blood under dark field microscopy. The vaccinated have blinking lights, rouleux formation, and what appears to be biocircuitry. The Pfizer trials warned the vaccinated cohorts to stay away from pregnant woman, warning that the vaccines can spread through skin contact and by aerosol. This shedding has been confirmed in Ana Mihalcea, MD, Ph.D's clinic. She looks at all her patients blood, both vaccinated and unvaccinated under electron microscopy and has found all the unvaccinated are showing the same things in the blood as the vaccinated. This is why Biden pushed so hard to reach a 70%+ vaccination rate. The goal all along was for this to shed onto the unvaccinated.

Go to her substack and you will find pictures of what's happening to the blood.

Here's an article that explains from the literature how the self assembling nanoparticles harvest your energy to build and run biocircuits inside of you.
https://anamihalceamdphd.su...

The Project Defuse, Peter Daszak and Ralph Baric's work was designed to create pathogens to scare people into taking these experimental vaccines. Go to howbadismybatch.com and note how many different lots there are and how they are identified. This is a mass experiment. There are many different types of these shots being tried out on people with an end goal of refining transhuman technologies.

4 0 Reply Share

Ektor
12 days ago

Three years too late. Nothing will change and they will continue to add more poison vaccines and mRNA substances to the childhood schedule in order to murder more children. That is the end game along with more profits for big pharma and the scumbag doctors who worship them.

3 0 Reply Share

Lionheart → Ektor
12 days ago

There has to be a way fight back and restore us back to the natural order of peace and excellent health.!

0 0 Reply Share

Stuff Happens
13 days ago

"If you get vaccinated you won't catch covid."
-Joe Biden

3 0 Reply Share

Jeanne DiMassa
13 days ago

Good luck with anything coming of this

4 1 Reply Share

IRISH
12 days ago

BAN all vaccines. ALL.

2 0 Reply Share

Afshin Nejat
12 days ago

Words we must never forget:
"Flatten the curve"
"Two more weeks"
"Over the nose!"
"Sorry, sir/mam, but masks are required"
"You're just being selfish"
"Think about grandma!"
"Just got the shot (wide-eyed sarcasm with mask on), haven't grown 3rd eye/turned into zombie, etc yet."
"They should be rounded up and put in quarantine"
"Mmm. I love these fries.... Mmmm"
and of course
"We're all in this together!"

These people are another species as far as I'm concerned.
General Blanchon just didn't go far enough in describing the disgracefulness of these (too many) people. But at least someone is trying to give credit where it is due. See "General Christian Blanchon" (French General)

2 0 Reply Share

Voyt Regnal → Afshin Nejat
7 days ago

..and: "do your part" 😉

1 0 Reply Share

r2b2judge
12 days ago

"The serious adverse events are all well documented," Mead said. "Yet it's surprising to see so many in the medical field continue to ignore or dismiss outright the latter half of the equation when considering all cause mortality trends."

Doctors are being threatened with losing their hard earned medical career if they speak out. Look at what happened to Dr.s Malik and Nass. What doctors want to be put through that for speaking up? It is not so surprising that doctors ignore or dismiss the Covid shots as being the cause of so many adverse events.

2 0 Reply Share

Len Hughes
7 days ago

It astounds me why these mRNA shots and the technology are being used and promoted. To say it's criminal is an understatement, it's outright evil at its core.

The harms are indescribable, those responsible need to face their accountability for the loss of millions of lives and ruination going forward of millions more through ruined health outcomes from these shots. Never did I think I'd be watching a slow marching genocide in my lifetime. Appears the Holocaust and Holodomor were typical for evil people to perpetrate on mankind. Will we EVER LEARN FROM THE PAST ON THIS????? Wish I could be positive, but I'm not so sure about that.

1 0 Reply Share

AL MCKNIGHT
7 days ago

It's way past time for the madness to end. Vaccines do more harm than good!

0 0 Reply Share

Miles

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